

- 22. The process of claim 20 or claim 21, wherein the phosphorylation or dephosphorylation is detected by using an intrinsic property of NDPK.
- 23. The process of claim 20 or claim 21, wherein the NDPK is modified to carry a label which gives a different detectable signal when the enzyme is phosphorylated from when it is unphosphorylated.
 - 24. The process of claim 23, wherein the NDPK carries a fluorescent label.
- The process of claim 24, wherein the fluorescent label is attached to the NDPK via a cysteine residue.
- 26. The process of claim 24, wherein the fluorescent label is IDCC (N-[2-(iodoacetamido)ethyl]-7-diethylaminocoumarin-3-carboxamide).
 - 27. The process of claim 20, wherein the nucleoside diphosphate is ADP or GDP.
 - 28. The process of claim 21, wherein the nucleoside triphosphate is ATP or GTP.
 - 29. The process of claim 20 or claim 21, being a quantitative process.

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30. The process of claim 20 or claim 21, wherein the NDPK is the NDPK of Myxococcus xanthus carrying a Asp112-Cys mutation, and carrying an IDCC label at this mutated residue.

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31. NDPK is modified to carry a label which gives a different detectable signal when the enzyme is phosphorylated from when it is unphosphorylated.

- 32. The NDPK of claim 31, wherein the label on the modified NDPK is a fluorescent label.
- 33. The NDPK of claim 32, wherein the fluorescent label is attached to the NDPK via a cysteine residue.
 - 34. The NDPK of claim 32 or claim 33, wherein the fluorescent label is IDCC.
- 35. NDPK of Myxococcus xanthus carrying a Aspl 12-Cys mutation, and carrying an IDCC label at this mutated residue.
- 36. NDPK modified by the attachment of at least one detectable label that is sensitive to the binding of a nucleoside diphosphate.
- 37. A substrate having the NDPK of any one of claims 31, 35 or 36 immobilised thereto.
- 38. The NDPK of any one of claims 31, 35 or 36 for use as an *in vivo* or *in vitro* diagnostic reagent.